BPA-ICD9 Diagnosis Code Listing

Alphabetical Order

WA Register of Developmental Anomalies

Description	Code	Major
17 ALPHA HYDROXYLASE DEF SYNTH	25524	Y
21 HYDROXYLASE DEF SYNTHESIS	25520	Y
47,XXX	75885	Y
47,XYY	75884	Y
ABNORMAL LIMB UNSPECIFIED	75548	Y
ABSENCE DIGITS UNSPECIFIED	75544	Y
ABSENCE OF BRAIN	74000	Y
ABSENT NIPPLE	75763	N
ACCESSORY AURICLE/S	74410	N
ACCESSORY DIGIT/S UNSPECIFIED	75509	N
ACCESSORY FINGER/S	75500	N
ACCESSORY HAND/FOREARM	75504	Y
ACCESSORY THUMB/S	75501	N
ACCESSORY TOE/S	75502	N
ACHILLES TENDON/S SHORT	75472	Y
ACHONDROPLASIA	75643	Y
ACRANIA	74001	Y
ADDISONS DISEASE	25540	Y
ADENOMA LIVER	21150	Y
ADENOMYOMA GASTRIC PYLORIS	21100	Y
ADRENAL ANOMALY OTHER	75918	Y
ADRENAL GLAND ABSENT	75910	Y
ADRENAL GLAND ANOMALY UNSPEC	75919	Y
ADRENAL GLAND ECTOPIC	75913	N
ADRENAL HYPERPL CONG NO SALT L	25526	Y
ADRENAL HYPERPL CONG SALT LOSS	25525	Y
ADRENAL HYPOPLASIA	75911	Y
ADRENOGENTIAL DIS OTHER&UNSPEC	25529	Y
ADRENOLEUCODYSTROPHY	33030	Y
AFIBRINOGENAEMIA	28630	Y
AGAMMAGLOBULINAE CONG SEX LINK	27900	Y
AGAMMAGLOBULINAEMIA X-LINKED	27901	Y
ALBINISM	27020	Y
ALIMENTARY TRACT ANOM SPEC	75080	Y
ALPHA 1 ANTITRYPSIN DEFICIENCY	27762	Y
ALVEOLAR NOTCH/CLEFT	74914	N
AMINO-ACID BRNCH/CHN DISTURBAN	27039	Y
AMINO-ACID METAB DISTURBANCES	27029	Y
AMINO-ACID TRNSPT DISTRB OTHER	27009	Y
AMNIOTIC BAND SYNDROME	76280	Y
ANAL ATRESIA/IMPERF NO FISTULA	75124	Y
ANAL ATRESIA/IMPERF WITH FISTU	75123	Y
ANAL FISTULA	75154	Y
ANAL STENOSIS	75125	Y
ANENCEPHALY	74002	Y
ANGIOMA/ANGIOFIBROMA/LIPOMA	22900	Y
ANKLE ANOMALIES	75562	Y
ANODONTIA/OLIGO/ABSENT	52000	N
ANOPHTHALMOS	74300	Y
ANTERIOR EYE ANOMALY OTHER	74348	Y
ANUS ECTOPIC/ANTERIORLY PLACED	75153	Y
ANUS/INTEST/CYST DUPLICATION	75150	Y
AORTA ANOMALY OTHER SPEC	74728	Y
AORTA ANOMALY UNSPECIFIED	74729	Y
AORTA OVERRIDING	74726	Y
AORTIC ARCH INTERRUPTED	74712	Y
AORTIC ARCH R SIDED	74723	Y
AORTIC ATRESIA	74720	Y
AORTIC HYPOPLASIA	74721	Y
AORTIC ROOT DILATATION	74727	Y
AORTIC STENOSIS SUPRA	74722	Y
AORTIC VALVE BICUSPID	74640	Y
AORTIC VALVE DYSPLASIA	74631	Y
AORTIC VALVE STENOSIS	74630	Y
AORTOPULMONARY WINDOW	74501	Y
APHAKIA	74330	Y
ARGINOSUCCINIC ACID DEFICIENCY	27060	Y
ARM ANOMALIES OTHER SPECIFIED	75556	Y
ARM/S ABSENT (HAND PRESENT)	75521	Y
ARM/S ABSENT (NO HAND)	75520	Y
ARM/SHOULDER ANOM OTHER SPEC	75558	Y
ARM/SHOULDER ANOM UNSPECIFIED	75559	N
ARNOLD CHIARI MALF NO SPIN BIF	74227	Y
ARTERIOVENOUS ANEURYSM BRAIN	74780	Y
ARTERIOVENOUS MALF PERIPHERAL	74762	Y
ARTHRITIS CONGENITAL	71690	Y
ARTHROGRYPOSIS	75580	Y
ASD NOS	74559	Y
ASD SPECIFIED	74558	Y
ASPHYXIATING THORACIC DYS	75640	Y
ASPLENIA	75900	Y
ATRIOVENTRICULAR SEPTAL DEFECT	74562	Y
ATRIUM &/OR VENTRICLE DILATED	74677	Y
AUDITORY CANAL ABSENT	74400	Y
AURICLE ABSENT	74401	Y
AUTOSOMAL ANOMALY UNSPECIFIED	75859	Y
AUTOSOMAL DELETION SYN OTHER	75838	Y
BARTTER SYNDROME	25511	Y
BATTEN'S DISEASE	36271	Y
BECKER MUSCULAR DYSTROPHY	35911	Y
BILIARY ATRESIA	75165	Y
BILIARY TRACT ANOMALY NOS	75168	Y
BLACKFAN DIAMOND SYN/ANAEMIA	28402	Y
BLADDER AND/OR URETHRA ABSENT	75380	Y
BLADDER ANOMALY UNSPECIFIED	75392	Y
BLADDER DIVERTICULUM	75382	Y
BLADDER ECTOPIC	75381	Y
BLADDER EXSTROPHY	75350	Y

Description	Code	Major
BLADDER NECK OBSTR/ATR/STEN	75361	Y
BLADDER/URETHRA ANOM OTHER	75388	Y
BODY ASYMMETRY	75582	Y
BODY WALL ANOMALY	75679	Y
BONE DEMINERALISATION	27590	Y
BRACHYSYNDACTYLY FINGERS	75514	Y
BRACHYSYNDACTYLY TOES	75515	Y
BRAIN ANOMALIES OTHER SPEC	74248	Y
BRAIN ANOMALY NOS	74290	Y
BRAIN ANOMALY OTHER SPECIFIED	74228	Y
BRAIN ANOMALY UNSPECIFIED	74229	Y
BRAIN/SPINAL CORD ANOM OTHER	74259	Y
BRANCHIAL ACH/CYS/FST/RMNT/SNS	74440	N
BRANCHIAL ARCH ANOM OTHER	74448	N
BRON/TRAC/LARY ANOM UNSPEC	74839	Y
BRON/TRACH/LARY ANOM OTHER	74838	Y
BRONCHIAL ANOMALY	74835	Y
BRONCHIAL STENOSIS	74834	Y
BRONCHO-OESOPHAGEAL FISTULA	75033	Y
BUTTERFLY VERTEBRAE	75615	N
CAFFEY DISEASE	75653	Y
CARBAMOYL PHOS SYNT DEFICIENCY	27061	Y
CARBOHYDRATE METAB DISORD NOS	27190	Y
CARNITINE TRANSPORTER DEFECT	27288	Y
CARTILAGE ANOMALY NOS	75693	N
CATARACT	74332	Y
CAUDA EQUINA ANOMALY	74253	Y
CCAM	74848	Y
CENTRAL CORE MYOPATHY	35900	Y
CEREBELLAR ANOMALY	74223	Y
CEREBRAL CYST	74243	Y
CEREBRAL CYSTS MULTIPLE	74242	Y
CEREBRAL DEGENERATIVE DIS	33080	Y
CEREBRAL INFARCTION	43490	Y
CEREBRAL VESSEL ANOMALY	74781	Y
CEREBRUM ANOMALY	74220	Y
CERVIX ABSENT	75240	Y
CHARCOT-MARIE-TOOTH	35610	Y
CHEDIAK-HIGASHI SYNDROME	28820	Y
CHERUBISM	52680	Y
CHEST WALL ANOM OTHER	75482	Y
CHOANAL ATRESIA	74800	Y
CHOANAL STENOSIS/NARROWING	74801	Y
CHOLEDOCHAL CYST	75166	Y
CHOLESTEATOMA CONGENITAL	38530	Y
CHONDRODYSPLASIA	75641	Y
CHONDRODYSPLASIA & HAEMANGIOMA	75642	Y
CHORDEE	75262	Y
CHOROID ANOMALY	74353	Y
CHR 13 DELETION LONG ARM	75833	Y
CHR 17 OR 18 DELETION LONG ARM	75834	Y
CHR 17 OR 18 DELETION SHORT AR	75835	Y
CHROMOSOMAL MICRODELETION	75836	Y
CHROMOSOMAL MICRODUPLICATION	75856	Y
CHROMOSOME ADDITIONAL MARKER	75855	Y
CHROMOSOME ANOMALY OTHER	75858	Y
CHROMOSOME UNSPEC OTHER &UNSPE	75899	Y
CIRCULATING ENZYMES DEF OTHER	27769	Y
CIRCULATORY SYSTEM OTHER SPEC	74788	Y
CIRRHOSIS INDIAN CHILDHOOD	57152	Y
CIRRHOSIS OTHER & UNSPECIFIED	57159	Y
CITRULLINAEMIA	27069	Y
CLAW FOOT	75471	Y
CLEFT HARD PALATE BILAT	74901	Y
CLEFT HARD PALATE CENTRAL	74902	Y
CLEFT HARD PALATE NOS	74903	Y
CLEFT HARD PALATE UNILAT	74900	Y
CLEFT LIP BILAT	74911	Y
CLEFT LIP BILAT & CLEFT PALATE	74921	Y
CLEFT LIP CENTR & CLEFT PALATE	74922	Y
CLEFT LIP CENTRAL	74912	Y
CLEFT LIP INC UNI & ALV NOTCH	74916	Y
CLEFT LIP INCOMP & CLEFT PALAT	74927	Y
CLEFT LIP INCOMPLETE	74913	Y
CLEFT LIP NOS	74919	Y
CLEFT LIP NOS & CLEFT PALATE	74929	Y
CLEFT LIP UNI & ALV NCH & CL P	74924	Y
CLEFT LIP UNI & ALVEOLAR NOTCH	74915	Y
CLEFT LIP UNILAT	74910	Y
CLEFT LIP UNILAT & CLEFT PALAT	74920	Y
CLEFT PALATE INCOMPLETE NOS	74909	Y
CLEFT SOFT PALATE BILAT	74905	Y
CLEFT SOFT PALATE CENTRAL	74906	Y
CLEFT SOFT PALATE NOS	74907	Y
CLEFT SOFT PALATE UNILAT	74904	Y
CLITORAL ANOMALY	75245	Y
CLOACAL ANOMALY	75155	Y
CLOTTING DEFIC OTHER & NOS	28639	Y
CLUB HAND/S	75484	Y
CLUBFOOT UNSPECIFIED	75473	Y
CMV CONGENITAL	77110	Y
COAGULATION DEF OTHER	28690	Y
COARCTATION AORTA NOS	74719	Y
COARCTATION AORTA OTHER	74718	Y
COARCTATION AORTA POSTDUCTAL	74711	Y
COARCTATION AORTA PREDUCTAL	74710	Y
COELIAC DISEASE	57900	Y
COGAN'S SYNDROME	74381	Y
COLLODION BABY SYNDROME	75711	Y
COLOBOMA EYELID	74368	Y
COLOBOMA IRIS	74343	Y
COLOBOMA LENS	74334	Y
COMMON (SINGLE) VENTRICLE	74530	Y
COMMON AV CANAL DEFECT	74563	Y
CONG ALOPECIA	75740	N
CONGENITAL ANOMALY UNSPECIFIED	75999	Y

Description	Code	Major	Description	Code	Major
CONGENITAL BANDS	75142	Y	DUANE ANOMALY/SYNDROME	37870	Y
CONGENITAL HEART BLOCK	74687	Y	DUCHENNE MUSCULAR DYSTROPHY	35910	Y
CONJOINED TWINS HEAD	75941	Y	DUCT ARTERIOSUS PREM CLOSURE	74701	Y
CONJOINED TWINS OTHER	75948	Y	DUODENAL ATRESIA	75110	Y
CONJOINED TWINS THORAX	75942	Y	DUODENAL WEB	75156	Y
CONJOINED TWINS DICEPHALUS	75940	Y	DUPLICATION OF CHROMOSOME NOS	75893	Y
CONJOINED TWINS ISCHIOPAGUS	75945	Y	DWARF SYNDROMES OTHER	75644	Y
CONJOINED TWINS STER/PELVIS	75943	Y	DYSAUTONOMIA CONGENITAL	74281	Y
CONJOINED TWINS UNSPECIFIED	75949	Y	DYSERYTHROPOIETIC ANAEMIA	28581	Y
CONNECTIVE/MUSC ANOM NOS	75690	N	DYSMORPHIC/UNUSUAL FACIES	74491	N
CONNECTIVE/MUSC ANOM OTHER	75688	Y	EAR/S ANOMALY NOS	74430	N
CONNECTIVE/MUSCLE/TENDON ANOM	75588	Y	EAR/S ANOMALY OTHER	74428	N
CORNEAL ANOMALY	74341	Y	EAR/S MISPLACED	74424	N
CORNEAL DYSTROPHY	37150	Y	EAR/S MISSHAPEN	74423	N
CORNEAL OPACITY	74340	Y	EBSTEIN'S ANOMALY	74620	Y
CORPUS CALLOSUM ANOMALY	74221	Y	ECTODERMAL DYSPLASIA	75734	Y
CORTICOADRENAL INSUFF UNSPEC	25549	Y	ECTOPIA CORDIS	74682	Y
CRANIORACHISCHISIS	74010	Y	ECTROPION	74361	N
CRANIOSYNOSTOSIS	75600	Y	EHLERS-DANLOS SYNDROME 11	75685	Y
CRI-DU-CHAT SYNDROME	75831	Y	EISENMENGER SYNDROME	74541	Y
CRIGLER-NAJJAR TYPE SYNDROME	27740	Y	ELBOW/S DISLOCATED	75483	Y
CROUZON SYNDROME	75601	Y	ELBOW/UPPER ARM ANOMALIES	75554	Y
CUTIS LAXA HYPERELASTICA	75737	Y	ELLIPTOCYTOSIS HEREDITARY	28210	Y
CYSTIC FIBROSIS	27700	Y	ELLIS-VAN CREVELD SYNDROME	75652	Y
CYSTIC HYGROMA	75964	Y	EMPHYSEMA CONGENITAL	77029	Y
CYSTINOSIS	27000	Y	ENCEPHALOCOELE OCCIPITAL	74200	Y
CYSTINURIA	27001	Y	ENCEPHALOCOELE OTHER SPECIFIED	74208	Y
DANDY WALKER MALFORMATION	74231	Y	ENCEPHALOCOELE UNSPECIFIED	74209	Y
DDH BILATERAL	75430	Y	ENCEPHALOCOELE/AC MALF/HC/OCCI	74201	Y
DDH CONFIRMED NFI	75434	Y	ENCEPHALOCOELE/HC/OTHER SITES	74203	Y
DDH L	75431	Y	ENDOCARDIAL CUSHION DEF NOS	74569	Y
DDH R	75432	Y	ENDOCRINE GLAND ANOMALY OTHER	75928	Y
DDH UNILATERAL UNSPECIFIED	75433	Y	ENTROPION	74362	N
DDH UNSPECIFIED	75439	Y	ENZYME DEF ANAEMIA OTHER	28230	Y
DENTAL ABNORMALITIES	52020	N	EPIDERMOLYSIS BULLOSA	75733	Y
DENTAL MALFORMATIONS	52040	N	EPIGLOTTIS ANOMALY SPECIFIED	75081	Y
DENTAL MALOCCLUSION	52400	Y	EPILUS MAXILLA CONGENITAL	52380	Y
DENTINOGENESIS IMPERFECTA	52050	Y	EPIPHYSEAL DYSPLASIA MULTIPLE	75657	Y
DERMOID CYST	75739	N	EPISPADIAS	75261	Y
DEXTROCARDIA WITH SITUS INVERS	75930	Y	ERYTHROPOIT PROTOPOPHYRIA	27712	Y
DEXTROCARDIA WITHOUT SITUS INV	74680	Y	EUSTACHIAN TUBE ABSENT	74425	Y
DIABETES INSIPIDUS NEPHROGENIC	58810	Y	EVENTRATION OF DIAPHRAGM	75662	Y
DIABETES INSIPIDUS NOS	25351	Y	EXENCEPHALY	74008	Y
DIAPHRAGM ABSENT	75660	Y	EXOMPHALOS	75670	Y
DIAPHRAGMATIC ANOMALY OTHER	75668	Y	EXOPHTHALMOS	74388	Y
DIAPHRAGMATIC HERNIA	75661	Y	EXOSTOSIS MULTIPLE	75647	Y
DIAPHYSEAL DYPLASIA PROGRESS	75655	Y	EYE ANOMALY NOS	74390	Y
DIASTEMATOMYELIA	74252	Y	EYE ANOMALY OTHER	74380	Y
DIGESTIVE SYSTEM ANOMALY	75188	Y	EYELID ANOMALY OTHER	74363	N
DISACCHARIDASE DEFICIENCY	27136	Y	FACIAL ASYMMETRY	75400	Y
DOUBLE IN/OUTLET L VENTRICLE	74674	Y	FACIAL CLEFT	74928	Y
DOUBLE IN/OUTLET R VENTRICLE	74673	Y	FACIOSCAPULOHUMORAL DYSTROPHY	35912	Y
DOWN SYNDROME NOS	75809	Y	FACTOR 11 [XI] DEFICIENCY	28620	Y

Description	Code	Major			
FALLOPIAN TUBE ABSENT	75210	Y	GYRAL ANOMALY	74225	Y
FALLOPIAN TUBE ANOMALY	75219	Y	HAEMANGIOMA	75738	N
FAMILIAL ACTH (GLUCOID DEF)	25548	Y	HAEMANGIOMA IN ORGAN	22800	N
FANCONI ANAEMIA	28401	Y	HAEMOCHROMATOSIS NEONATAL	27502	Y
FANCONI SYNDROME	27002	Y	HAEMOGLOBIN H DISEASE	28274	Y
FAVS/GOLDENHAR SYNDROME	75606	Y	HAEMOGLOBINOPATHY OTHER	28279	Y
FEMALE GENITALIA ANOM UNSPEC	75249	Y	HAEMOLYTIC ANAEMIA CONG	28290	Y
FEMUR BOWED	75440	Y	HAEMOLYTIC URAEMIC SYNDROME	28310	Y
FETAL ALCOHOL SYNDROME/FASD	75992	Y	HAEMOPHAGOCYTIC LYMPHOHISTIOCY	20290	Y
FETUS ANATOMICAL DISRUPTION	75993	Y	HAEMOPHILIA A FACTOR 8 [VIII]	28600	Y
FIBROSARCOMA CONGENITAL	17100	Y	HAEMOPHILIA B FACTOR 9 [IX]	28610	Y
FINGER ANOMALIES	75550	N	HAEMORRAGHIC TELANGIECTASI	44800	Y
FINGER/S FUSED	75510	Y	HAIR ANOMALY NOS	75791	N
FISTULA-URETH/UTERUS/RECT	75232	Y	HAMARTOMA	75968	Y
FOOT ANOMALIES	75561	Y	HAMARTOMA NOS	75969	Y
FOOT ANOMALIES OTHER	75478	Y	HAND ANOMALIES	75551	Y
FOREARM & HAND ABSENT	75523	Y	HAND/S & OR FINGER/S ABSENT	75524	Y
FOREARM ABSENT (HAND PRESENT)	75522	Y	HARTNUP DISEASE	27003	Y
FOREARM ANOMALIES	75553	Y	HEARING LOSS	74287	Y
FRAGILE X	75888	Y	HEART ANOMALY CONG UNSPEC	74699	Y
FRUCTOSE INTOLERANCE HEREDITAR	27120	Y	HEART ANOMALY CYANOTIC NOS	74693	Y
FUSED VERTEBRAE	75616	N	HEART ANOMALY OTHER SPEC	74688	Y
G6PD DEFICIENCY	28220	Y	HEART HYPERPLASIA (DILATED)	74675	Y
GALACT-1-PHOS URIDYL TRANS DEF	27110	Y	HEMIPARESIS CONGENITAL	34310	Y
GALACTOKINASE DEFICIENCY	27112	Y	HEMIVERTEBRAE	75614	N
GALACTOSAEMIA OTHER & UNSPEC	27119	Y	HEPATIC/BILE DUCTS ANOMALY	75167	Y
GALLBLADDER ABSENT/AGENESIS	75163	Y	HERPES SIMPLEX CONGENITAL	77122	Y
GALLBLADDER ANOMALY OTHER	75164	Y	HETEROTOPIA MALFORMATION BRAIN	74245	Y
GANGLIONEUROMA	22500	Y	HIP ANOMALIES	75566	Y
GASTROSCHISIS	75671	Y	HIRSCHSPRUNG'S DIS LONG & SHOR	75130	Y
GENITALIA AMBIGUOUS FEMALE	75248	Y	HIRSCHSPRUNG'S DIS LONG SEGMN	75131	Y
GENITALIA AMBIGUOUS MALE	75288	Y	HIRSCHSPRUNG'S DIS SHORT SEGMN	75132	Y
GENU RECURVATUM	75443	Y	HIRSCHSPRUNG'S DISEASE NOS	75133	Y
GERBODE DEFECT	74542	Y	HISTIOCYTOSIS	27786	Y
GIGANTISM	25300	Y	HOLOPROSENCEPHALY	74226	Y
GLAUCOMA	74320	Y	HOMOCYSTINURIA	27040	Y
GLIOMA NASAL	19500	Y	HORNER SYNDROME	33790	Y
GLIOMA OPTIC NERVE	19051	Y	HYDRANENCEPHALY	74232	Y
GLOSSOPTOSIS	75013	N	HYDROCEPH ABSENT SEPTUM PEL	74234	Y
GLUTARIC ANOMALY	27089	Y	HYDROCEPHALUS DUE TO CYSTS	74233	Y
GLYCOGENOSIS OTHER & UNSPEC	27109	Y	HYDROCEPHALUS OTHER SPEC	74238	Y
GLYCOGENOSIS TYPE 1	27100	Y	HYDROCEPHALUS UNSPECIFIED	74239	Y
GLYCOGENOSIS TYPE 2 (POMPE'S)	27101	Y	HYDROCEPHALUS-AQUEDUCT STEN	74230	Y
GLYCOGENOSIS TYPE 3	27102	Y	HYDROMYELIA SPINAL CORD	74254	Y
GLYCOGENOSIS TYPE 4	27103	Y	HYDRONEPHROSIS	75320	Y
GLYCOSYLATION DISORDER CONG	27189	Y	HYDRONEPHROSIS A/N ONLY	75322	Y
GONADAL DYSGENESIS OTHER	75868	Y	HYDRONEPHROSIS A/N, NFI	75323	Y
GORLIN SYNDROME	17300	Y	HYDROPS FETALIS/ASCITES	77800	Y
GRANULOMATOUS DISEASE	28810	Y	HYPERBILIRUBINAEMIA CONG	27744	Y
GREAT TOE/S DUPLICATION	75503	N	HYPERCHOLESTEROLAEMIA FAMILIAL	27200	Y
GREAT VEIN ANOMALY NOS	74749	Y	HYPERCHYLOMICRONAEMIA	27230	Y
GREAT VEIN ANOMALY OTHER	74748	Y	HYPERGLYCINAEMIA	27070	Y
GROWTH HORMONE DEFICIENCY	25330	Y	HYPERINSULINAEMIA	25110	Y

Description	Code	Major	Description	Code	Major
HYPERTELORISM	75602	N	JEJUNAL ATRESIA	75111	Y
HYPERTYROSINAEMIA	27021	Y	KARTAGENER SYNDROME	75934	Y
HYPOGAMMAGLOBULINANAEMIA	27904	Y	KASABACH-MERRIT SYNDROME	28738	Y
HYPOPARATHYROIDISM	25210	Y	KIDNEY ACCESSORY	75330	Y
HYPOPHOSPHATASIA OTHER SPEC	27532	Y	KIDNEY ANOMALY SPECIFIED	75338	Y
HYPOPHOSPHATASIA RICKETS	27531	Y	KIDNEY ANOMALY UNSPECIFIED	75339	Y
HYPOPLASTIC ATRIUM	74678	Y	KIDNEY ANOMALY UNSPECIFIED	75390	Y
HYPOPLASTIC L HEART SYNDROME	74670	Y	KIDNEY CYSTIC DISEASE OTHER	75318	Y
HYPOPLASTIC L VENTRICLE	74672	Y	KIDNEY CYSTIC DISEASE UNSPEC	75319	Y
HYPOPLASTIC R HEART	74671	Y	KIDNEY DUPLEX	75331	Y
HYPOPLASTIC R VENTRICLE	74676	Y	KIDNEY DYSPLASIA NO CYSTS BIL	75304	Y
HYPOSPADIAS CONFIRMED NFI	75266	Y	KIDNEY DYSPLASIA NO CYSTS UNIL	75305	Y
HYPOSPADIAS CONFIRMED NOS SITE	75269	Y	KIDNEY ECTOPIC	75333	Y
HYPOSPADIAS CORONAL	75264	Y	KIDNEY HORSESHOE	75332	Y
HYPOSPADIAS GLANULAR	75263	Y	KIDNEY HYPOPLASIA BILATERAL	75302	Y
HYPOSPADIAS MIDSHAFT	75267	Y	KIDNEY HYPOPLASIA UNILATERAL	75303	Y
HYPOSPADIAS NOS	75260	Y	KIDNEY MEDULLARY CYSTIC JUV	75314	Y
HYPOSPADIAS PENOSCROTAL	75265	Y	KIDNEY MULTICYSTIC DYSPLASIA	75316	Y
HYPOSPADIAS PERINEAL	75268	Y	KIDNEY MULTICYSTIC NO DYSPLA	75317	Y
HYPOTHALAMUS ANOMALY	74222	Y	KIDNEY/S HYPERPLASTIC	75334	Y
HYPOTHYROIDISM CONG	24399	Y	KIDNEY/S POLYCYSTIC OTHER NOS	75313	Y
ICHTHYOSIS CONG (HARLEQUIN)	75710	Y	KIDNEY/S POLYCYSTIC ADULT TYPE	75312	Y
ICHTHYOSIS CONGENITA OTHER	75719	Y	KIDNEY/S POLYCYSTIC INFANTILE	75311	Y
IGA DEFICIENCY	27902	Y	KLINEFELTER PHENO EXTRA CHROM	75871	Y
IGG IMMUNO DEFICIENCY	27909	Y	KLINEFELTERS 47XXY	75870	Y
ILEAL ATRESIA	75112	Y	KLINEFELTERS NOS	75879	Y
IMMUNODEFICIENCY NOS	27930	Y	KLIPPEL-FEIL SYNDROME	75611	Y
IMMUNODEFICIENCY X-LINKED	27980	Y	KNEE ANOMALIES	75564	Y
INCONTINENTIA PIGMENTI	75735	Y	KNEE DISLOCATION	75444	Y
INDETERMINATE SEX NOS	75279	Y	KOILONYCHIA CONGENITAL	75752	N
INFANTILE SPASMS X-LINKED	34580	Y	KYPHOSCOLIOSIS NO VERT ANOMALY	75424	Y
INFERIOR VENA CAVA STENOSIS	74740	Y	KYPHOSCOLIOSIS WITH VERT ANOM	75612	Y
INIENCEPHALY CLOSED	74020	Y	KYPHOSIS NO VT ANOM-GIVE SITE	75423	Y
INIENCEPHALY OPEN	74021	Y	LACRIMAL ANOMALY	74366	N
INIENCEPHALY UNSPECIFIED	74029	Y	LACTIC ACIDOSIS	27622	Y
INNER EAR ANOMALY	74403	Y	LACTOSE INTOLERANCE CONG	27130	Y
INSENSITIVITY TO PAIN CONG	74282	Y	LARGE BOWEL ATRESIA/BLND END	75120	Y
INTEGUMENT OTHER SPECIFIED	75780	Y	LARSEN SYNDROME	75581	Y
INTERSEX 46XX	75272	Y	LARYNGEAL WEBBING	74820	Y
INTERSEX 46XY	75271	Y	LARYNX ANOMALY	74830	Y
INTERSEX OTHER	75276	Y	LEBER'S AMAUROSIS	36272	Y
INTERSEX TRUE GONADAL	75270	Y	LEG/HIP/PELVIS ANOM OTHER SPEC	75568	N
INTERSEX UNSPECIFIED	75273	y	LEG/HIP/PELVIS ANOM UNSPEC	75569	Y
INTESTINAL ANOMALY OTHER	75158	Y	LEG/S ABSENT (FOOT PRESENT)	74531	Y
INTESTINAL ANOMALY UNSPECIFIED	75159	Y	LEG/S ABSENT (FOOT PRESENT)	75531	Y
INTRA UTERINE INFECT NOS	77189	Y	LEG/S ABSENT (NO FOOT)	75530	Y
INTRACRANIAL HAEMORRHAGE	43200	Y	LENS ANOMALY OTHER	74339	Y
INTRAORAL DEFORMITY MAJOR	74484	Y	LENS ECTOPIC	74333	Y
IRIS ABSENT	74342	Y	LETTERER-SIWE DISEASE	20250	Y
IRIS ANOMALY OTHER	74344	Y	LEUCODYSTROPHY/KRABBE DISEASE	33000	Y
IRON METABOLISM DISORDER	27509	Y	LEUKAEMIA CONGENITAL NOS	20890	Y
IVEMARK SYNDROME	75906	Y	LEUKAEMIA LYMPHOBLASTIC CONG	20400	Y
JAW ANOMALY	75408	Y	LEUKAEMIA MONOCYTIC ACUTE CONG	20600	Y

Description	Code	Major			
LEUKAEMIA MYELOID CONG	20500	Y	LYMPHOEDEMA CONGENITAL	75700	Y
LIMB ANOMALY UNSPEC	75590	Y	MACROGLOSSIA	75012	N
LIMB ASYMMETRY	75583	Y	MACROPHTHALMIA	74321	Y
LIMB REDUCTION DEF UNSPECIFIED	75549	Y	MACROSTOMIA	74480	N
LIP ANOMALY OTHER	75027	N	MACROTIA	74420	N
LIP PITS/SUB-MUCOUS	75026	N	MACULAR DYSTROPHY	36250	Y
LIPODYSTROPHY	27260	Y	MAJOR VESSEL/ARTERY ABNORMAL	74737	Y
LIPOMA	21400	Y	MALABSORPTION CARBOH MONOSACCH	57981	Y
LIPOMENING/AC MALF/HC/CV	74153	Y	MALABSORPTION UNSPECIFIED	57989	Y
LIPOMENING/AC MALF/HC/LM	74155	Y	MALIG NEOPLASM BRAIN	19100	Y
LIPOMENING/AC MALF/HC/LM-SC	74156	Y	MALIG NEOPLASM CONNECTIVE TISS	17190	Y
LIPOMENING/AC MALF/HC/SC	74157	Y	MALIG NEOPLASM ENDOCRINE GLAND	19400	Y
LIPOMENING/AC MALF/HC/SITE NOS	74159	Y	MALROTATION CAECUM & COLON	75140	Y
LIPOMENING/AC MALF/HC/TH	74154	Y	MALROTATION OTHER & UNSPEC	75149	Y
LIPOMENING/AC MALF/NO HC/CV	74163	Y	MARCUS GUNN JAW-WINK SYN	74280	Y
LIPOMENING/AC MALF/NO HC/LM	74165	Y	MATERNAL DRUG INDUCED ANOMALY	75990	Y
LIPOMENING/AC MALF/NO HC/LM-SC	74166	Y	MCAD DEFICIENCY	27283	Y
LIPOMENING/AC MALF/NO HC/NOS	74169	Y	MCCUNE ALBRIGHT SYNDROME	75651	Y
LIPOMENING/AC MALF/NO HC/SC	74167	Y	MECKEL-GRUBER SYNDROME	75970	Y
LIPOMENING/AC MALF/NO HC/TH	74164	Y	MECKELS DIVERTICULUM	75101	Y
LIPOMENING/HC/CV	74133	Y	MEGACOLON CONGENITAL	75134	Y
LIPOMENING/HC/LM	74135	Y	MEGALENCEPHALY	74240	Y
LIPOMENING/HC/LM-SC	74136	Y	MEGALO CORNEA	74322	Y
LIPOMENING/HC/SC	74137	Y	MENINGOCOELE CRANIAL	74204	Y
LIPOMENING/HC/SITE NOS	74139	Y	MENINGOCOELE OTHER SITES	74205	Y
LIPOMENING/HC/TH	74134	Y	MESENTERIC REMNANT CYST	75211	Y
LIPOMENING/NO HC/CV	74143	Y	MESENTERY ANOMALY	75141	Y
LIPOMENING/NO HC/LM	74145	Y	MESOBLASTIC NEPHROMA	23690	Y
LIPOMENING/NO HC/LM-SC	74146	Y	METABOLIC ACIDAEMIA	27621	Y
LIPOMENING/NO HC/SC	74147	Y	METABOLIC ACIDOSIS NOS	27623	Y
LIPOMENING/NO HC/SITE NOS	74149	Y	METABOLIC DISORDER CONG UNSPEC	27790	Y
LIPOMENING/NO HC/TH	74144	Y	METABOLIC DISORDER CONG UNSPEC	74299	Y
LISSENCEPHALY/AGYRIA	74224	Y	METABOLIC DISORDER OTHER SPEC	27788	Y
LIVER ABSENT/AGENESIS	75160	Y	METABOLIC DISTURBANCES SYN	75987	Y
LIVER ANOMALY OTHER	75162	Y	METABOLIC KETOACIDAEMIA	27626	Y
LIVER CYSTIC DISEASE	75161	Y	METAPHYSEAL DYSPLASIA	75645	Y
LONGIT REDUCT DEF UNSPECIFIED	75543	Y	METATARSUS VARUS	75452	Y
LONGIT REDUCT DEF UPPER LIMBS	75527	Y	METHYLMALONIC ACIDEMIA/URIA	27079	Y
LONGT RED DEF LEG/FOOT/TOES	75536	Y	MICRO PENIS	75285	Y
LORDOSIS NO VT ANOM-GIVE SITE	75421	Y	MICROCEPHALY	74210	Y
LOWER LEG & FOOT ABSENT	75533	Y	MICROCOLON	75152	Y
LOWER LEG ABSENT (FOOT PRESENT	75532	Y	MICROGASTRIA	75070	Y
LOWER LIMB ANOMALIES	75563	Y	MICROGNATHIA	75409	N
LUNG ABSENT	74850	Y	MICROPHTHALMOS	74310	Y
LUNG ACCESSORY LOBE	74862	N	MICROSTOMIA	74481	N
LUNG ANOMALY OTHER & NOS	74869	Y	MICROTIA	74421	Y
LUNG CYST SINGLE	74840	Y	MIDDLE EAR ANOMALY	74402	Y
LUNG CYSTS MULTIPLE	74841	Y	MIDFACIAL HYPOPLASIA	74488	Y
LUNG DYSPLASIA OTHER & NOS	74858	Y	MITOCHONDRIAL ABNORMALITY	75994	Y
LUNG INTERSTITIAL DISEASE	74868	Y	MITRAL LEAFLET SINGLE	74652	Y
LUNG SEQUESTRATION	74852	Y	MITRAL VALVE ANOMALY	74653	Y
LUNG/S HYPOPLASIA	74851	Y	MITRAL VALVE ANOMALY NOS	74655	Y
LYMPHATIC ABNORMALITY OTHER	75965	Y	MITRAL VALVE ATRESIA	74654	Y

Description	Code	Major
MITRAL VALVE CLEFT	74651	Y
MITRAL VALVE INCOMPETENCE	74660	Y
MITRAL VALVE LEAFLET ANOMALY	74656	Y
MITRAL VALVE STENOSIS	74650	Y
MOSAIC INCLUDING XXXXY	75883	Y
MOTOR NEURONE DISEASE	33520	Y
MOTORSENSORY NEUROPATH HERED	35620	Y
MOUTH/PHARYNX ANOM OTHER	75028	Y
MOYAMOYA SYNDROME	43750	Y
MUCOLIPIDOSES	27270	Y
MUCOPOLYSACCHARIDOSIS 1 SYN	27751	Y
MUCOPOLYSACCHARIDOSIS 2 SYN	27752	Y
MUCOPOLYSACCHARIDOSIS 3 SYN	27753	Y
MUCOPOLYSACCHARIDOSIS 4 SYN	27754	Y
MUCOPOLYSACCHARIDOSIS 6 SYN	27756	y
MUCOPOLYSACCHARIDOSIS 7 SYN	27757	Y
MULLERIAN DUCT CYST PERSISTENT	75287	Y
MULT ENDOCRINE NEOPLASIA	23740	Y
MULTIPLE SKELETAL ABNORM	75699	Y
MUSCLE ABSENT	75681	Y
MUSCULAR DYSTROPHY CONG NOS	35909	Y
MUSCULAR DYSTROPHY OTHER&UNSPE	35919	Y
MUSCULOSKELETAL OTHER SPEC	75488	N
MYASTHENIA GRAVIS CONGENITAL	35809	Y
MYELOFIBROMA	21500	Y
MYOCARDIAL ANOMALY	74686	Y
MYOFIBROMA TONGUE	21000	Y
MYOFIBROMATOSIS	23810	Y
MYOPATHY OTHER	35980	Y
MYOPATHY CONGENITAL NOS	35903	Y
MYOPATHY MULTICORE CONGENITAL	35908	Y
MYOPATHY UNSPECIFIED	35990	Y
MYOTONIC DYSTROPHY	35920	Y
MYOTUBULAR MYOPATHY	35901	Y
NAIL ANOMALIES NOS	75792	N
NAIL ANOMALY OTHER	75758	N
NAIL DYSPLASIA	75755	N
NAIL HYPOPLASIA	75750	N
NAIL-PATELLA SYNDROME	75683	Y
NASAL AGENESIS/RUDIMENTARY	74810	Y
NASAL ANOMALY OTHER SPEC	74818	Y
NASAL ANOMALY UNSPECIFIED	74819	Y
NASAL CLEFT DEFORMITY	74812	Y
NECK WEBBED	74450	N
NEMALINE ROD MYOPATHY	35902	Y
NEPHROPATHY HEREDITARY	58324	Y
NEPHROTIC SYNDROME CONG	75337	Y
NERVOUS SYSTEM ANOMALIES SPEC	74288	Y
NEUROBLASTOMA	19410	Y
NEURODEGENERATIVE DISOR UNSPEC	33090	Y
NEUROFIBROMATOSIS	23770	Y
NEUTROPENIA CONGENITAL	28800	Y
NEUTROPENIA CYCLICAL	28804	Y
NYSTAGMUS	37950	Y
OCULAR MOTILITY DISORDER	37880	Y
OCULOMOTOR APRAXIA	31540	Y
ODONTOGENIC CYSTS	52600	Y
OESOPHAGEAL ATRESIA NO FISTULA	75030	Y
OESOPHAGEAL DUPLICATION	75043	Y
OESOPHAGEAL STRICTURE/STENOSIS	75034	Y
OESOPHAGUS ANOMALY OTHER	75048	Y
OLIGOHYDRAMNIOS SEQUENCE	76120	Y
OMPHALOMESENTERIC DUCT PERSIS	75100	Y
OPTIC DISC ANOMALY	74352	Y
ORBIT ANOMALY	74367	Y
ORCHIOBLASTOMA	18690	Y
OSEOPH/TRACHEAL ANOM OTHER	75038	Y
OSTEOCHONDROMATA	21300	Y
OSTEODYSTROPHIES UNSPEC	75659	Y
OSTEOGENESIS IMPERFECTA	75650	Y
OSTEOPETROSIS	75654	Y
OSTEOPOROSIS CONGENITAL	73300	Y
OSTIUM PRIMUM DEFECT	74560	Y
OSTIUM SECUNDUM DEFECT	74551	Y
OVARY/IES ABSENT	75200	Y
OVARY/IES ANOMALY OTHER	75208	Y
OVARY/IES ANOMALY OTHER	75209	Y
OVARY/IES STREAK	75201	Y
PACHYONYCHIA CONGENITA	75751	N
PALATE ANOMALY OTHER	75025	N
PANCREAS ABSENT/HYPOPLASIA	75170	Y
PANCREAS ACCESSORY	75171	Y
PANCREAS ANNULAR	75172	Y
PANCREAS ECTOPIC	75173	Y
PANCREATIC ANOMALY NOS	75179	Y
PANCREATIC ANOMALY OTHER	75178	Y
PANCREATIC EXOCRIN INSUFF	57780	Y
PANHYPOPITUITARISM	25329	Y
PARATHYROID ANOMALY	75923	Y
PARTIAL ANOM PUL VEIN RET/DRAN	74743	Y
PARTIAL AV CANAL DEFECT	74568	Y
PDA	74700	Y
PECTUS CARINATUM	75480	N
PECTUS EXCAVATUM	75481	N
PELVIS ANOMALIES	75567	Y
PELVI-URETERIC JNCT OBST	75321	Y
PENDRED SYNDROME	24390	Y
PENILE ANOMALY OTHER	75286	Y
PENTALOGY OF CANTRELL	75678	Y
PENTALOGY OF FALLOT	74521	Y
PERICARDIAL ANOMALY	74685	Y
PERIPHERAL ARTERY ANOMALY	74764	Y
PERIPHERAL VASC ANOMALY OTHER	74768	Y
PERIPHERAL VASCULAR ANOM NOS	74769	Y
PERIPHERAL VEIN ANOMALY	74765	Y
PERSIS L SVC TO COR SINUS	74741	Y

Description	Code	Major
PES CAVUS	75470	Y
PES PLANUS VALGUS	75461	Y
PEUTZ-JEGHER SYNDROME	75960	Y
PHARYNGEAL ANOMALY	75021	Y
PHIMOSIS CONGENITAL	60500	Y
PHLEBECTASIA CONG	74763	y
PHOCOMELIA UNSPECIFIED	75541	Y
PIERRE ROBIN SYNDROME	75603	Y
PILI TORTI	75742	N
PINEAL TUMOUR NOS	23700	Y
PITUITARY ANOMALY	75920	Y
PITUITARY DWARFISM OTHER	25331	Y
PITUITARY HORMONE DEFICIENCY	25342	Y
PKU	27010	Y
PLATELET DYSFUNCTION	28710	Y
POLAND ANOMALY	75680	Y
PORENCEPHALIC CYST	74241	Y
PORTAL VEIN-HEPATIC ARTERY FIS	74745	Y
POST SEGMENT ANOM OTHER	74359	Y
POSTERIOR URETHRAL VALVES	75360	Y
POTTER'S FACIES	75401	N
POTTERS SYNDROME	75306	Y
PRE-AURICULAR PIT/SINUS/FISTUL	74441	N
PRECOCIOUS PUBERTY	25910	Y
PRIMARY DENTITION ANOMALY	52060	Y
PROGEROID SYNDROME	25980	Y
PRUNE-BELLY SYNDROME	75672	Y
PSEUDO HYPOALDOSTERONISM	25543	Y
PSEUDOHYPOPARATHYROIDISM	27542	Y
PTOSIS	74360	N
PULM ART ATRESIA & SEPTAL DEF	74731	Y
PULM ARTERY ANOM OTHER	74738	Y
PULM ARTERY ANOMALY NOS	74739	Y
PULMONARY ARTERY ATRESIA	74730	Y
PULMONARY ARTERY DILATATION	74733	Y
PULMONARY ARTERY STENOSIS	74732	Y
PULMONARY STENOSIS INFUNDIBULA	74683	Y
PULMONARY TELANGIECTASES	74843	Y
PULMONARY VALVE ANOM OTHER	74608	Y
PULMONARY VALVE ANOM UNSPEC	74609	Y
PULMONARY VALVE ATRESIA	74600	Y
PULMONARY VALVE ATRESIA (IVS)	74603	Y
PULMONARY VALVE INCOMPETENCE	74602	Y
PULMONARY VALVE STENOSIS	74601	Y
PYKNOCYTOSIS HEREDITARY	28391	Y
PYLORIC OBSTRUCT OTHER	75058	Y
PYLORIC STENOSIS	75051	Y
PYRUVATE DEHYDROG DEFICIENCY	27185	Y
RADIAL APLASIA	75526	Y
RANULA	75022	N
RECTAL ATRES/ABSENT NO FISTULA	75122	Y
RECTAL ATRESIA WITH FISTULA	75121	Y
RECTO-URETHAL FISTULA	75386	Y
RECTO-VAGINAL FISTULA	75242	Y
REDUCTION DEF LOWER LIMB OTHER	75538	Y
REDUCTION DEF LOWER LIMB UNSPE	75539	Y
REDUCTION DEF UPPER LIMB OTHER	75528	Y
REDUCTION DEF UPPER LIMB UNSPE	75529	Y
REFSUM'S DISEASE	35630	Y
RENAL AGENESIS BILATERAL	75300	Y
RENAL AGENESIS UNILATERAL	75301	Y
RENAL ARTERY ANOM OTHER	74761	Y
RENAL ARTERY STENOSIS	74760	Y
RENAL CYST SINGLE	75310	Y
RENAL PELVIS ANOMALY OTHER	75328	Y
RENAL PELVIS ANOMALY UNSPEC	75329	Y
RENAL PELVIS BIFID	75336	Y
RENAL SYSTEM ANOM UNSPECIFIED	75399	Y
RESPIRATORY CHAIN DEFICIENCY	27780	Y
RESPIRATORY SYS ANOM OTHER	74888	Y
RETINAL ANOMALY	74351	Y
RETINITIS PIGMENTOSA	36270	Y
RETINOBLASTOMA	19050	Y
RHABDOMYOMA HEART	21270	Y
RHABDOMYOMA NON-CARDIAC	21590	Y
RHABDOMYOSARCOMA BLADDER	18900	Y
RHABDOMYOSARCOMA LUNG	16200	Y
RIB ANOMALY OTHER SPECIFIED	75634	N
RIB/S ABSENT	75630	N
RIB/S CERVICAL	75620	N
RIB/S EXTRA	75633	N
RIB/S MISSHAPEN	75631	N
RIBS FUSED	75632	N
RICKETS VIT D RESISTANT	27530	Y
RICKETTS CONG	26800	Y
ROGERS DISEASE	74540	Y
RUBELLA CONGENITAL	77100	Y
SACRAL VERTEBRAE ANOMALY	75617	N
SALIVARY GLAND ANOMALY	75023	Y
SCHEUERMANNS DISEASE	73200	Y
SCOLIOSIS NO VT ANOM-GIVE SITE	75420	Y
SCROTAL & TESTIS ANOM OTHER	75282	Y
SEPTAL DEFECTS OTHER SPEC	74580	Y
SEPTAL DEFECTS UNSPECIFIED	74590	Y
SHORT STATURE	75692	N
SHORTENED UPPER & OR LOWER ARM	75525	Y
SHORTENED UPPER & OR LOWER LEG	75535	Y
SHORTENED UPPER&LOWER LIMBS	75983	Y
SHOULDER ANOMALIES	75555	N
SICKLE CELL ANAEMIA	28260	Y
SIDEROBLASTIC ANAEMIA	28501	Y
SINGLE COMMON ATRIUM	74561	Y
SITUS INVERSUS ABDOMINIS	75933	Y
SITUS INVERSUS AMBIGUOUS	75935	Y
SITUS INVERSUS NOS	75939	Y
SITUS INVERSUS THORACIS	75932	Y

Description	Code	Major	Description	Code	Major
SITUS INVERSUS WITH LEVOCARDIA	75931	Y	SPLEEN MISSHAPEN	75903	N
SKELETAL DYSPLASIA OTHER	75658	Y	SPONDYLOEPIPHYSEAL DYSPLA	75646	Y
SKELETAL SYNDROME N.E.C.	75986	Y	SPONDYLOLISTHESIS	75613	Y
SKIN ANOMALY NOS	75790	Y	STERNUM ABSENT	75635	Y
SKIN ANOMALY SYND OTHER	75730	Y	STERNUM ANOMALY OTHER	75638	N
SKULL ANOMALY OTHER SPECIFIED	75608	N	STEROID SULPHATASE DEFICIENCY	27758	Y
SKULL ASYMMETRY	75405	Y	STOMACH ANOMALY OTHER	75078	Y
SKULL DEFECTS LOCALISED	75607	N	STOMACH DIVERTICULUM	75074	Y
SKULL&FACE BONES ANOM UNSPEC	75609	N	STOMACH DUPLICATION	75075	Y
SMALL BOWEL ATRESIA	75119	Y	STURGE-WEBER SYNDROME	75961	Y
SOTOS SYN/CEREBRAL GIGANTISM	25981	Y	SUBAORTIC ANEURYSM	74724	Y
SPACES BETW INCISORS	52080	N	SUBGLOTTIC ATRESIA/STENOSIS	74831	Y
SPADE-LIKE HAND/S	75485	N	SUCROSE INTOLERANCE	27134	Y
SPBIF/AC MALF/HC/CV	74113	Y	SUCROSE-ISOMALTASE INTOLERANCE	27135	Y
SPBIF/AC MALF/HC/LM	74115	Y	SYNDACTYLY FINGERS	75511	N
SPBIF/AC MALF/HC/LM-SC	74116	Y	SYNDACTYLY TOES	75513	N
SPBIF/AC MALF/HC/SC	74117	Y	SYNDACTYLY UNSPECIFIED	75519	N
SPBIF/AC MALF/HC/SITE NOS	74119	Y	SYNDROME INVOLVING LIMBS	75984	Y
SPBIF/AC MALF/HC/TH	74114	Y	SYNDROME OF FACE N.E.C.	75980	Y
SPBIF/AC MALF/NO HC/CV	74123	Y	SYNDROME SHORT STATURE	75982	Y
SPBIF/AC MALF/NO HC/LM	74125	Y	SYNDROME UNCLASSIFIED	75988	Y
SPBIF/AC MALF/NO HC/LM-SC	74126	Y	SYNDROMES OTHER SPECIFIED	75989	Y
SPBIF/AC MALF/NO HC/SC	74127	Y	SYNOPHRYS	75748	N
SPBIF/AC MALF/NO HC/TH	74124	Y	SYPHILIS CONGENITAL	09090	Y
SPBIF/ACMALF/NO HC/SITE NOS	74129	Y	SYRINGOMYELIA	33600	Y
SPBIF/HC/CV	74103	Y	TALIPES CALCANEOVALGUS	75460	Y
SPBIF/HC/LM	74105	Y	TALIPES CALCANEOVARUS	75451	Y
SPBIF/HC/LM-SC	74106	Y	TALIPES EQUINOVARUS BILAT	75454	y
SPBIF/HC/SC	74107	Y	TALIPES EQUINOVARUS L	75456	y
SPBIF/HC/SITE NOS	74109	Y	TALIPES EQUINOVARUS R	75455	Y
SPBIF/HC/TH	74104	Y	TALIPES EQUINOVARUS UNSPEC	75450	Y
SPBIF/NO HC/CV	74193	Y	TAR SYNDROME	28732	Y
SPBIF/NO HC/LM	74195	Y	TAY-SACHS DISEASE	33010	Y
SPBIF/NO HC/LM-SC	74196	Y	TERATOMA CERVICAL	23880	Y
SPBIF/NO HC/SC	74197	Y	TERATOMA INTERCRANIAL/CEREBRAL	23750	Y
SPBIF/NO HC/SITE NOS	74199	Y	TERATOMA INTRAABDOMINAL	23500	Y
SPBIF/NOHC/TH	74194	Y	TERATOMA MEDIASTINAL	23580	Y
SPBIF/OP/NO HC/CV-SC	74192	Y	TERATOMA OROFACIAL	23510	Y
SPHERICAL LENS	74331	Y	TERATOMA SACROCOCCYGEAL	23800	Y
SPHEROCYTOSIS HEREDITARY	28200	Y	TERATOMA TESTIS	22200	Y
SPINAL CORD ANOMALY	74258	Y	TESTICULAR HYPOPLASIA	75281	Y
SPINAL CORD HYPOPL/DYSPLASIA	74251	Y	TESTICULAR TORSION CONGENITAL	75258	Y
SPINAL MUSCLE HYPOPLASIA	35880	Y	TESTIS ABSENT	75280	Y
SPINAL MUSCULAR ATROPHY	33510	Y	TESTIS ECTOPIC	75253	Y
SPINOCEREBELLAR ATAXIA	33430	Y	TETRALOGY OF FALLOT	74520	Y
SPINOCEREBELLAR OTHER SPEC	33480	Y	TETRALOGY OF FALLOT VARIANT	74522	Y
SPINOCEREBELLAR UNSPEC	33490	Y	TGA COMPLETE	74510	Y
SPLEEN ACCESSORY	75904	Y	TGA CORRECTED	74512	Y
SPLEEN ANOMALY SPECIFIED	75908	Y	TGA INCOMPLETE	74511	Y
SPLEEN ANOMALY UNSPECIFIED	75909	Y	TGA NOS	74519	Y
SPLEEN ECTOPIC	75905	Y	TGA OTHER SPECIFIED	74518	Y
SPLEEN HYPERPLASIA	75902	Y	THALASSAEMIA ALPHA	28240	Y
SPLEEN HYPOPLASIA	75901	Y	THALASSAEMIA BETA INTERMEDIA	28243	Y

Description	Code	Major
THALASSAEMIA BETA MAJOR	28244	Y
THALASSAEMIA MAJOR OTHER	28248	Y
THORACIC CAGE ANOMALY	75639	Y
THROMBOCYTOPAE AMEGAKARYOCYTIC	28739	Y
THROMBOCYTOPENIA	28731	Y
THROMBOPHILIA	28679	Y
THYMUS ANOMALY	75924	Y
THYROGLOSSAL CYST/REM/FISTULA	75922	N
THYROID ANOMALY	75921	Y
THYROID DYSHORMONGENESIS	24610	Y
TIBIA/FIBULA BOWING	75441	Y
TOE/S AND OR FEET ABSENT	75534	Y
TOE/S ANOMALY	75560	N
TOE/S FUSED	75512	Y
TOF WITH OESOPHAGEAL ATRESIA	75031	Y
TOF WITHOUT OESOPH ATRESIA	75032	Y
TONGUE ABSENT	75010	Y
TONGUE ANOMALY OTHER	75018	Y
TONGUE BIFID/GROOV/CLFT/FORKED	75014	N
TONGUE HYPOPLASIA	75011	Y
TOTAL ANOM PUL VEIN RET/DRAIN	74742	Y
TOXOPLASMOSIS CONGENITAL	77121	Y
TRACHEAL ATRESIA/AGENESIS	74833	Y
TRANSLOCATIONS OTHER	75854	Y
TRANSPOSITION BOWEL & APPENDIX	75151	Y
TREACHER COLLINS SYNDROME	75604	Y
TRICUSPID VALVE ANOM OTHER/NOS	74613	Y
TRICUSPID VALVE ATRESIA	74610	Y
TRICUSPID VALVE MALALIGNED	74614	Y
TRICUSPID VALVE OVERRIDING	74618	Y
TRICUSPID VALVE REGURGITATION	74612	Y
TRILOGY OF FALLOT	74684	Y
TRIPLOIDY	75857	Y
TRISCUPID VALVE CLEFT	74611	Y
TRISOMY 13	75810	Y
TRISOMY 13 MOSAIC	75814	Y
TRISOMY 13 TRANSLOCATION	75812	Y
TRISOMY 18	75820	Y
TRISOMY 18 TRANSLOCATION	75822	Y
TRISOMY 21	75800	Y
TRISOMY 21 DUPLICAT, TRANSLOCA	75802	Y
TRISOMY 21 MOSAIC	75804	Y
TRISOMY 21 OTHER SPECIFIED	75805	Y
TRISOMY 8	75850	Y
TRISOMY OTHER	75851	Y
TRISOMY PARTIAL	75853	Y
TRISOMY TOTAL OTHER	75852	Y
TROYER SYNDROME	33410	Y
TRUNCUS ARTERIOSUS	74500	Y
TUBEROUS SCLEROSIS	75950	Y
TURNERS KARYOTYPE NORMAL	75862	Y
TURNERS KARYOTYPE UNSPEC	75869	Y
TURNERS MOSAIC	75861	Y
TURNERS SYNDROME	75860	Y
UDT BILAT	75251	Y
UDT INDETERMINATE	75254	Y
UDT NOS	75252	Y
UDT UNILAT	75250	Y
UDT VANISHING/REGRESSION	75257	Y
UMBILICAL CORD ABNORMALITY	76268	Y
UPPER LEG ANOMALIES	75565	Y
URACHAL REMNANT	75370	N
URETER ABSENT	75340	Y
URETER ANOMALY SPECIFIED	75348	Y
URETER DUPLEX	75341	Y
URETER ECTOPIC	75342	Y
URETER/S DILATED	75345	Y
URETERIC ANOMALY UNSPECIFIED	75391	Y
URETHRA &/OR MEATUS DOUBLE	75384	Y
URETHRA ANT OBSTR/ATRESIA/STEN	75362	Y
URETHRA ECTOPIC	75385	Y
URETHRAL ANOMALY UNSPECIFIED	75393	Y
URETHRAL FISTULA NOS	75387	Y
URETHRAL OBSTRUCT OTHER&UNSPEC	75369	Y
URINARY MEATUS OBSTR/ATR/STEN	75363	Y
URTICARIA PIGMENTOSA	75732	Y
UTERINE AGENESIS	75230	Y
UTERINE ANOMALY OTHER	75238	Y
UTERUS BIFID	75220	Y
UVULA BIFID/CLEFT/DUPLIC/BILOB	74908	N
VAGINA ABSENT	75241	Y
VAGINAL CYST	75247	Y
VALGUS DEFORMITY UNSPECIFIED	75469	Y
VARICELLA ZOSTER CONGENITAL	77129	Y
VARUS DEFORMITIES UNSPECIFIED	75459	Y
VARUS DEFORMITY COMPLEX	75453	Y
VAS & PROSTATE ANOM OTHER	75284	Y
VAS DEFERENS ABSENT	75283	Y
VASCULAR MALFORMATION BRAIN	74244	Y
VASCULAR RING /DOUBLE AORTA	74725	Y
VENTRICULOMEGALY	74235	Y
VERTEBRAE ANOMALY OTHER SPECIF	75618	N
VERTEBRAE/SPINE ANOMALY NOS	75619	N
VESICO-URETERIC JUNCT OBS	75343	Y
VESICO-URETERIC REFLUX	75344	Y
VIRAL INFECTION CONGENITAL	77125	Y
VITREOUS HUMOUR ANOMALY	74350	Y
VON HIPPEL LINDAU SYNDROM	75962	Y
VON WILLEBRAND DISEASE	28640	Y
VSD MUSCULAR	74548	Y
VSD NOS	74549	Y
VULVA ABSENT OR OTHER ANOMALY	75244	Y
WERDNIG-HOFFMAN DISEASE	33500	Y
WISKOTT-ALDRICH SYNDROME	27911	Y
WOLFF-HIRSCHORN SYNDROME	75832	Y
WRIST ANOMALIES	75552	Y

Description	Code	Major
XANTHOGRANULOMA JUVENILE	27220	N
XO/XX MOSAIC	75881	Y
XO/XXY MOSAIC	75882	Y
XO/XY MOSAIC	75880	Y